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APPENDIX I

CLAIM AMENDMENTS:

Cancel Claims 5, 8, 17 and 20, and enter new Claims 23 to 26, as indicated in the following listing of the claims:

- 1. (previously presented) A choline ascorbate in form of crystals and having diffraction lines at d = 3.80 Å and 4.55 Å which are most intense in a range between 3.40 and 4.70 Å in a 2 Ø X-ray powder diffractogram.
- (previously presented) A choline ascorbate in form of crystals, wherein the crystals are free from water of crystallization.
- 3. (canceled)
- 4. (previously presented) The choline ascorbate crystals as claimed in claim 1, having an intensity ratio of the diffraction lines at d = 3.80 Å and d = 4.55 Å of at least 0.5.
- 5. (canceled)
- 6. (previously presented) A process for preparing choline ascorbate in form of crystals having diffraction lines at d = 3.80 Å and 4.55 Å which are most intense in a range between 3.40 and 4.70 Å in a 2 Θ X-ray powder diffractogram, which comprises reacting ascorbic acid with triethylamine and ethylene oxide, and carrying out the reaction in a temperature range from -10°C to 40°C.
- (previously presented) The process of claim 6, which is carried out in a water-miscible organic solvent.
- 8. and 9. (canceled)
- 10. (previously presented) Drugs comprising the choline ascorbate claimed in claim 1.
- 11. (previously presented) Additives in foods, additives in animal feeds or food supplements comprising the choline ascorbate claimed in claim 1.
- 12. (previously presented) The process of claim 6, wherein ascorbic acid is reacted with triethylamine and ethylene oxide by adding ethylene oxide to a mixture comprising the ascorbic acid and the triethylamine.

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- 13. (previously presented) The process of claim 12, wherein gaseous ethylene oxide is added to the mixture comprising the ascorbic acid and the triethylamine.
- 14. (previously presented) A choline ascorbate in form of anhydrous crystals having a melting point from 123.5 to 124.4°C or in the range from 123.5 to 124.4°C.
- 15. (previously presented) The choline ascorbate crystals as claimed in claim 2, having diffraction lines at d=3.80 Å and 4.55 Å which are most intense in a range between 3.40 and 4.70 Å in a 2 Θ X-ray powder diffractogram.
- 16. (previously presented) The choline ascorbate crystals as claimed in claim 2, having diffraction lines at d = 3.80 Å and d = 4.55 Å which have an intensity ratio of at least 0.5.
- 17. (canceled)
- 18. (previously presented) A process for preparing the choline ascorbate defined in claim 2, which comprises reacting ascorbic acid with triethylamine and ethylene oxide, and carrying out the reaction in a temperature range from -10°C to 40°C.
- 19. (previously presented) The process of claim 18, which is carried out in a water-miscible organic solvent.
- 20. (canceled)
- 21. (previously presented) A drug comprising the choline ascorbate crystals defined in claim 2.
- 22. (previously presented) An additive in foods or in animal feeds or a food supplement comprising the choline ascorbate crystals defined in claim 2.
- 23. (new) A process for preparing choline ascorbate, wherein the choline ascorbate is obtained in form of anhydrous crystals having diffraction lines at d = 3.80 Å and 4.55 Å which are most intense in a range between 3.40 and 4.70 Å in a 2 Θ X-ray powder diffractogram and having a melting point from 123.5 to 124.4°C or in the range from 123.5 to 124.4°C, which process comprises
 - a) providing a mixture of ascorbic acid, triethylamine and a solvent,
 - b) adding to the mixture gaseous ethylene oxide, and

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c) crystallizing the choline ascorbate,

wherein stages (a) and (b) are carried out at a temperature of from -10°C to 40°C, and

the solvent is a water miscible organic solvent or is a mixture of said organic solvent and water.

- 24. (new) The process of Claim 23, wherein the solvent is a water miscible organic solvent.
- 25. (new) The process of Claim 23, wherein the choline ascorbate is crystallized from the solvent employed in stage (a).
- 26. (new) The choline ascorbate obtained by the process of claim 23.

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